

**Response Under 37 CFR § 1.116 \* -- Expedited Procedure -- Examining Group 1631  
Docket No. 1073.060  
U.S. Serial No. 09/595,096**

**Amendments to the Claims:**

This listing of claims will replace all prior versions and listings, of claims in the application:

**Listing of Claims:**

1. **(Currently Amended)** A computer-aided method of ~~docking a ligand to~~ modeling ligand- protein having a binding site complexes so as to identify ligands likely to have therapeutic activity, said method comprising:

performing a pre-docking conformational search and generating multiple solution conformations of the a ligand therefrom;

generating a binding site image of the a protein, said binding site image comprising multiple hot spots;

matching hot spots of the binding site image to atoms in at least one conformation of the multiple solution conformations of the ligand to obtain at least one position of the ligand relative to the protein in a protein-ligand complex; and

optimizing the at least one position of the ligand while allowing translation, orientation and rotatable bonds of the ligand to vary, and while holding the protein fixed.

2. **(Previously Presented)** The method of claim 1, additionally comprising, after performing the pre-docking conformational search and generating multiple solution conformations, creating a database of the multiple solution conformations of the ligand and storing said three-dimensional database for subsequent use by said matching.

3. **(Previously Presented)** The method of claim 2, wherein said database of the multiple solution conformations of the ligand comprises a conformational database of a combinatorial library.

4. **(Currently Amended)** The method of claim 1, wherein said performing the pre-docking conformational search and generating multiple solution conformations of the ligand comprises:

randomly generating a plurality of conformations of the ligand;

**Response Under 37 CFR § 1.116 -- Expedited Procedure -- Examining Group 1631**  
**Docket No. 1073.060**  
**U.S. Serial No. 09/595,096**

minimizing ~~a~~the strain of each conformation of the plurality of conformations;  
using the strain and a the solvent accessible surface area of each conformation  
to rank the conformations; and  
clustering the conformations and retaining a desired top number of clusters of  
conformations, said retained top number of clusters of conformations comprising said  
multiple conformations of the ligand in solution.

5. **(Currently Amended)** The method of claim 1, wherein said generating the  
binding site image includes at least one of creating a list of ~~a~~polar apolar hot spots identifying  
points in the binding site that are favorable for an ~~a~~polar apolar atom to bind, and generating a  
list of polar hot spots identifying points in the binding site that are favorable for a hydrogen bond  
donor or acceptor to bind.

6. **(Previously Presented)** The method of claim 5, wherein said generating the  
binding site image further comprises:

placing a grid around the binding site of the protein;  
determining a hot spot search volume using said grid;  
determining hot spots using a grid-like search of the hot spot search volume; and  
for each type of hot spot, clustering the hot spots and retaining a desired number  
of top clusters of hot spots, said desired number of top clusters comprising said multiple  
hot spots to be employed by said matching.

7. **(Previously Presented)** The method of claim 1, wherein said matching  
comprises:

matching atoms of the at least one solution conformation of the ligand to  
appropriate hot spots of the protein by positioning the at least one solution conformation  
of the ligand as a rigid body into the binding site image;  
defining a match, said match determining a unique rigid body transformation; and  
using the unique rigid body transformation to place the at least one solution  
conformation of the ligand into the binding site of the protein.

8. **(Previously Presented)** The method of claim 7, wherein said determining the  
unique rigid body transformation comprises determining the unique rigid body transformation  
that minimizes:

*Response Under 37 CFR § 1.116 -- Expedited Procedure -- Examining Group 1631*  
*Docket No. 1073.060*  
*U.S. Serial No. 09/595,096*

$$I(R,T) = \sum_{j=1}^J |H_j - RA_j - T|^2$$

where:

$I(R,T)$  = rms deviation between a  $j^{th}$  hot spot and a  $j^{th}$  atom of the at least one solution conformation of the ligand;

$H_j$  = a position vector of a  $j^{th}$  hot spot of the protein;

$A_j$  = a position vector of a  $j^{th}$  atom of the at least one solution conformation of the ligand;

$R$  = a 3x3 rotation matrix; and

$T$  = a translation vector.

9. **(Previously Presented)** The method of claim 1, wherein multiple positions of the ligand are obtained, and said optimizing step comprises

eliminating each position of the ligand having a predetermined percentage of atoms with a steric clash;

ranking remaining positions of the ligand using an atom pairwise score with a desired atom score cutoff, said atom pairwise score comprising a hydrogen bonding potential score or a steric potential score;

after ranking, clustering the positions of the ligand and selecting a top number  $n$  of positions; and

optimizing each of the  $n$  positions, allowing the translation, orientation and rotatable bonds of the ligand to vary.

10. **(Previously Presented)** The method of claim 9, wherein said optimizing comprises optimizing each position of the  $n$  positions using a Broyden-Fletcher-Goldfarb-Shanno (BFGS) optimization algorithm with said atom pairwise score, allowing the translation, orientation and rotatable bonds of the ligand to vary.

11. **(Currently Amended)** A computer-aided system for docking ~~a modeling~~ ligand-~~protein to a protein having a binding-site complexes so as to identify ligands likely to have therapeutic activity~~, said system comprising:

means for performing a pre-docking conformational search and generating multiple solution conformations of the ~~a~~ ligand therefrom;

**Response Under 37 CFR § 1.116 -- Expedited Procedure -- Examining Group 1631  
Docket No. 1073.060  
U.S. Serial No. 09/595,096**

means for generating a binding site image of ~~the~~a protein, said binding site image comprising multiple hot spots;

means for matching hot spots of the binding site image to atoms in at least one solution conformation of the multiple conformations of the ligand to obtain at least one position of the ligand relative to the protein in a protein-ligand complex; and

means for optimizing the at least one position of the ligand while allowing translation, orientation and rotatable bonds of the ligand to vary, and while holding the protein fixed.

12. **(Previously Presented)** The system of claim 11, additionally comprising means for creating a database of the multiple solution conformations of the ligand and for storing said three-dimensional database for subsequent use by said matching, after performing the pre-docking conformational search and generating multiple solution conformations of the ligand.

13. **(Previously Presented)** The system of claim 12, wherein said database of the multiple solution conformations of the ligand comprises a conformational database of a combinatorial library.

14. **(Currently Amended)** The system of claim 11, wherein said means for performing the pre-docking conformational search and generating multiple solution conformations of the ligand comprises:

means for randomly generating a plurality of conformations of the ligand;

means for minimizing ~~a~~the strain of each conformation of the plurality of conformations;

means for using the strain and ~~a~~the solvent accessible surface area of each conformation to rank the conformations; and

means for clustering the conformations and retaining a desired top number of clusters of conformations, said retained top number of clusters of conformations comprising said multiple solution conformations of the ligand .

15. **(Original)** The system of claim 11, wherein said means for generating the binding site image includes at least one of means for creating a list of apolar hot spots identifying points in the binding site that are favorable for an apolar atom to bind, and means for generating a list of polar hot spots identifying points in the binding site that are favorable for a

**Response Under 37 CFR § 1.116 \* -- Expedited Procedure -- Examining Group 1631**  
**Docket No. 1073.060**  
**U.S. Serial No. 09/595,096**

hydrogen bond donor or acceptor to bind.

16. **(Previously Presented)** The system of claim 15, wherein said means for generating the binding site image further comprises:

- means for placing a grid around the binding site of the protein;
- means for determining a hot spot search volume using said grid;
- means for determining hot spots using a grid-like search of the hot spot search volume; and
- for each type of hot spot, means for clustering the hot spots and for retaining a desired number of top clusters of hot spots, said desired number of top clusters comprising said multiple hot spots to be employed by said matching.

17. **(Previously Presented)** The system of claim 11, wherein said means for matching comprises:

- means for matching atoms of the at least one solution conformation of the ligand to appropriate hot spots of the protein by positioning the at least one solution conformation of the ligand as a rigid body into the binding site image;
- means for defining a match, said match determining a unique rigid body transformation; and
- means for using the unique rigid body transformation to place the at least one solution conformation of the ligand into the binding site of the protein.

18. **(Previously Presented)** The system of claim 17, wherein said determining the unique rigid body transformation comprises determining the unique rigid body transformation that minimizes:

$$I(R,T) = \sum_{j=1}^n |H_j - RA_j - T|^2$$

where:

$I(R,T)$  = rms deviation between a  $j^{\text{th}}$  hot spot and a  $j^{\text{th}}$  atom of the at least one solution conformation of the ligand ;

$H_j$  = a position vector of a  $j^{\text{th}}$  hot spot of the protein;

$A_j$  = a position vector of a  $j^{\text{th}}$  atom of the at least one solution conformation of the ligand ;

$R$  = a 3x3 rotation matrix; and

**Response Under 37 CFR § 1.116 \* -- Expedited Procedure -- Examining Group 1631  
Docket No. 1073.060  
U.S. Serial No. 09/595,096**

T = a translation vector.

19. **(Previously Presented)** The system of claim 11, wherein multiple positions of the ligand are obtained, and said means for optimizing comprises :

means for eliminating each position of the ligand having a predetermined percentage of atoms with a steric clash;

means for ranking remaining positions of the ligand using an atom pairwise score with a desired atom score cutoff, said atom pairwise score comprising a hydrogen bonding potential score or a steric potential score;

after ranking, means for clustering the positions of the ligand and selecting a top number n of positions; and

means for optimizing each of the n positions, allowing the translation, orientation and rotatable bonds of the ligand to vary.

20. **(Previously Presented)** The system of claim 19, wherein said means for optimizing comprises means for optimizing each position of the n positions using a Broyden-Fletcher-Goldfarb-Shanno (BFGS) optimization algorithm with said atom pairwise score, allowing the translation, orientation and rotatable bonds of the ligand to vary.

21. **(Currently Amended)** At least one program storage device readable by a machine, tangibly embodying at least one program of instructions executable by the machine to perform a method of ~~docking a ligand to a modeling ligand~~ protein having a binding site complexes so as to identify ligands likely to have therapeutic activity, said method comprising:

performing a pre-docking conformational search and generating multiple solution conformations of the a ligand therefrom;

generating a binding site image of the a protein, said binding site image comprising multiple hot spots;

matching hot spots of the binding site image to atoms in at least one conformation of the multiple solution conformations of the ligand to obtain at least one position of the ligand relative to the protein in a protein-ligand complex; and

optimizing the at least one position while allowing translation, orientation and rotatable bonds of the ligand to vary, and while holding the protein fixed.

**Response Under 37 CFR § 1.116 \* -- Expedited Procedure -- Examining Group 1631**  
**Docket No. 1073.060**  
**U.S. Serial No. 09/595,096**

22. **(Previously Presented)** The at least one program storage device of claim 21, additionally comprising, after performing the pre-docking conformational search and generating multiple solution conformations of the ligand, creating a database of the multiple solution conformations of the ligand and storing said three-dimensional database for subsequent use by said matching.

23. **(Previously Presented)** The at least one program storage device of claim 22, wherein said database of the multiple solution conformations of the ligand comprises a conformational database of a combinatorial library.

24. **(Currently Amended)** The at least one program storage device of claim 21, wherein said performing the pre-docking conformational search and generating multiple solution conformations of the ligand comprises:

- randomly generating a plurality of conformations of the ligand;
- minimizing ~~a the strain and a solvent accessible surface area~~ of each conformation of the plurality of conformations;
- using the strain and the solvent accessible surface area of each conformation to rank the conformations; and
- clustering the conformations and retaining a desired top number of clusters of conformations, said retained top number of clusters of conformations comprising said multiple solution conformations of the ligand.

25. **(Original)** The at least one program storage device of claim 21, wherein said generating the binding site image includes at least one of creating a list of apolar hot spots identifying points in the binding site that are favorable for an apolar atom to bind, and generating a list of polar hot spots identifying points in the binding site that are favorable for a hydrogen bond donor or acceptor to bind.

26. **(Previously Presented)** The at least one program storage device of claim 25, wherein said generating the binding site image further comprises:

- placing a grid around the binding site of the protein;
- determining a hot spot search volume using said grid;
- determining hot spots using a grid-like search of the hot spot search volume; and
- for each type of hot spot, clustering the hot spots and retaining a desired number

**Response Under 37 CFR § 1.116 \* – Expedited Procedure – Examining Group 1631**  
**Docket No. 1073.060**  
**U.S. Serial No. 09/595,096**

of top clusters of hot spots, said desired number of top clusters comprising said multiple hot spots to be employed by said matching.

27. **(Previously Presented)** The at least one program storage device of claim 21, wherein said matching comprises:

matching atoms of the at least one solution conformation of the ligand to appropriate hot spots of the protein by positioning the at least one solution conformation of the ligand as a rigid body into the binding site image;

defining a match, said match determining a unique rigid body transformation; and using the unique rigid body transformation to place the at least one solution conformation of the ligand into the binding site of the protein.

28. **(Previously Presented)** The at least one program storage device of claim 27, wherein said determining the unique rigid body transformation comprises determining the unique rigid body transformation that minimizes:

$$I(R,T) = \sum_{j=1}^N |H_j - RA_j - T|^2$$

where:

$I(R,T)$  = rms deviation between a  $j^{\text{th}}$  hot spot and a  $j^{\text{th}}$  atom of the at least one solution conformation of the ligand ;

$H_j$  = a position vector of a  $j^{\text{th}}$  hot spot of the protein;

$A_j$  = a position vector of a  $j^{\text{th}}$  atom of the at least one solution conformation of the ligand;

$R$  = a 3x3 rotation matrix; and

$T$  = a translation vector.

29. **(Previously Presented)** The at least one program storage device of claim 21, wherein multiple positions of the ligand are obtained, and said optimizing step comprises :

eliminating each position of the ligand having a predetermined percentage of atoms with a steric clash;

ranking remaining positions of the ligand using an atom pairwise score with a desired atom score cutoff, said atom pairwise score comprising a hydrogen bonding potential score or a steric potential score;



**Responso Under 37 CFR § 1.116 \* -- Expedited Procedure -- Examining Group 1631  
Docket No. 1073.060  
U.S. Serial No. 09/595,096**

after ranking, clustering the positions of the ligand and selecting a top number n of positions; and

optimizing each of the n positions, allowing the translation, orientation and rotatable bonds of the ligand to vary.

30. **(Previously Presented)** The at least one program storage device of claim 29, wherein said optimizing comprises optimizing each position of the n positions using a Broyden-Fletcher-Goldfarb-Shanno (BFGS) optimization algorithm with said atom pairwise score, allowing the translation, orientation and rotatable bonds of the ligand to vary.